

F₂ 20. (Amended) The transgenic mouse according to Claim 1, wherein Flp recombinase activity is further regulated by a factor selected from the group consisting of chemical, developmental stage and temperature.

sub G1 } 51. (Twice Amended) The transgenic mouse according to claim 15, wherein said another transgene is a gene controlling differentiation of a cell or development of an organism selected from the group consisting of genes encoding adhesion molecules, cyclin kinase inhibitors, Wnt family members, Pax family members, F₃ Winged helix family members, Hox family members, cytokines, interleukins, growth/differentiation factors and their receptors, kinases, phosphatases, metabolic enzymes, and antigen receptors.

sub G2 } 52. (Amended) A transgenic mouse comprising a Flp recombinase transgene intergrated into the genome of the transgenic mouse, wherein the Flp recombinase transgene is expressed from a tissue specific or a developmental stage specific promoter in at least one cell of the transgenic mouse at a level sufficient to F₄ catalyze recombination between two FLP-recognition sequences in direct repeat orientation in said cell, wherein said recombination is detected by activation of a gene expressed from a ubiquitous promoter, wherein said gene produces a detectable product only when in recombined form.

F5 55. (Amended) The transgenic mouse of claim 52, wherein said detectable product is a histochemical marker encoded by said gene selected from the group consisting of alkaline phosphatase, β -galactosidase, chloramphenicol acetyltransferase, luciferase, green fluorescent protein and β -glucuronidase.

56. (Amended) The transgenic mouse of claim 52, wherein said detectable product is a transcript expressed from said gene in recombined form that is detectable by *in situ* hybridization.

57. (Amended) The transgenic mouse of claim 52, wherein said detectable product is a peptide tag encoded by said gene that is detectable by binding to a cognate binder.

Sub G3 } 59. (Amended) A method of mapping the developmental fate of a cell *in vivo* comprising:

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- (a) providing a transgenic mouse comprising a genome which contains a Flp recombinase transgene under control of a tissue-specific or developmental stage specific promoter and at least two FLP recognition sequences in direct orientation;
 - (b) expressing the Flp recombinase transgene at a level sufficient to catalyze site-specific recombination between said FLP recognition sequences in at least one cell; and

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(c) ~~detecting said recombination in said at least one cell by detecting activation of a gene expressed from a ubiquitous promoter, wherein said gene produces a detectable product only when in recombined form, and wherein said recombination is evidence of expression of said F1p transgene in said cell or a developmental precursor to said cell.~~

62. (Amended) The method of claim 59, wherein said detectable product is a histochemical marker encoded by said gene selected from the group consisting of alkaline phosphatase, β -galactosidase, chloramphenicol acetyltransferase, luciferase, green fluorescent protein and β -glucuronidase.

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63. (Amended) The method of claim 59, wherein said detectable product is a transcript expressed from said gene in recombined form that is detectable by *in situ* hybridization.

64. (Amended) The method of claim 59, wherein said detectable product is a peptide tag encoded by said gene that is detectable by binding to a cognate binder.